LISTING OF CLAIMS

Claim 1 (currently amended): A compound of the formula (I)

$$Ar^{1} \bigvee_{\mathbf{Q}} \mathbf{Y} \bigvee_{\mathbf{R}^{5}} \mathbf{R}^{6} \bigvee_{(I);}^{\mathbf{R}^{y}}$$

wherein:

 Ar^1 is an aromatic carbocycle chosen from phenyl, naphthyl, tetrahydronaphthyl, indanyl and indenyl, each Ar^1 is optionally substituted with one R^1 , and wherein Ar^1 is independently substituted with two R^2 groups and wherein one R^1 and one R^2 on adjacent ring atoms optionally form a 5- or 6-membered carbocyclic or heterocyclic ring:

 R^1 is halogen, NO₂, NH₂, J-N(R^a)-(CH₂)_m-, N(J)₂-(CH₂)_m-, NH₂C(O)-, J-N(R^a)-C(O)-, J-S(O)_m- N(R^a)-S(O)_m- or heterocycle (CH₂)_m- wherein the heterocyclic group is optionally substituted by C_{1.5}-alkyl;

Q is a N or CR^p;

Y is
$$> CR^pR^v$$
, $-CR^p = C(R^v)$, $-O$, $-N(R^x)$ or $> S(O)_m$;

wherein R^a , R^p , R^v , R^x and R^y are each independently hydrogen or C_{1-5} alkyl;

X is
$$-CH_2$$
-, $-N(\mathbf{R}^a)$ -, -O- or -S-;

W is Nor CH;

each m is independently 0,1 or 2;

J is chosen from C1-10 alkyl and carbocycle each optionally substituted by R^b;

R² is chosen from C1-6 alkyl, C3-7 cycloalkyl optionally substituted by C1-5 alkyl, C1-4 acyl, aroyl, C1-4 alkoxy, each being optionally partially or fully halogenated, halogen, C1-6 alkoxycarbonyl, carbocyclesulfonyl and -SO₂-CF₃;

each R³, R⁴ and R⁵ are independently chosen from hydrogen, C1-6 alkyl and halogen;

 \mathbf{R}^{6} is optionally attached at a position *ortho* or *meta* to the N atom of the indicated ring, and is chosen from

a bond, -O-, -O-(CH₂)₁₋₅-, >C(O), -NH-, -C(O)-NH-, -S-, C_{1-5} alkyl branched or unbranched, C_{2-5} alkenyl, C_{1-3} acyl, C_{1-3} alkyl(OH), heterocycle selected from morpholinyl, piperazinyl, piperidinyl, pyrrolidinyl and tetrahydrofuranyl, heteroaryl selected from pyridinyl, pyrimidinyl, pyrazinyl, pyridazinyl, pyrrolyl, imidazolyl, pyrazolyl, thienyl, furyl, isoxazolyl, thiazolyl, oxazolyl and isothiazolyl or-aryl each alkyl, alkenyl, acyl , heterocycle, heteroaryl and aryl are optionally substituted by one to three hydroxy, oxo, C_{1-3} alkyl, C_{1-3} alkoxy, C_{1-5} alkoxycarbonyl, -NR₇R₈ or NR₇R₈-C(O)-:

wherein each R_6 is further optionally covalently attached to groups chosen from:

hydrogen, -N $\mathbf{R}_7\mathbf{R}_8$, C₁₋₃ alkyl, C₃₋₆ cycloalkylC₀₋₂alkyl, hydroxy, C₁₋₃ alkoxy, phenoxy, benzyloxy, arylC₀₋₄ alkyl, heteroaryl C₀₋₄alkyl and heterocycle C₀₋₄alkyl, each above listed heterocycle, heteroaryl and aryl group is optionally substituted by one to three hydroxy, oxo, C₁₋₄ alkyl, C₁₋₃ alkoxy, C₁₋₅ alkoxycarbonyl, N $\mathbf{R}_7\mathbf{R}_8$ -C(O)- or C₁₋₄ acyl;

each \mathbf{R}_7 and \mathbf{R}_8 are independently hydrogen, phenyl $C_{0\text{-}3}$ alkyl optionally subtituted by halogen, $C_{1\text{-}3}$ alkyl or di $C_{1\text{-}5}$ alkyl amino, or \mathbf{R}_7 and \mathbf{R}_8 are $C_{1\text{-}2}$ acyl, benzoyl or $C_{1\text{-}5}$ branched or unbranched alkyl optionally substituted by $C_{1\text{-}4}$ alkoxy, hydroxy or mono or di $C_{1\text{-}3}$ alkyl amino;

and

R^b is chosen from hydrogen, C1-5 alkyl, hydroxyC1-5 alkyl, C2-5 alkenyl, C2-5 alkynyl, carbocycle, heterocycle, heterocycle, heterocycle, C1-5 alkoxy, C1-5 alkylthio, amino, C1-5 alkylamino, C1-5 dialkylamino, C1-5 acyl, C1-5 alkoxycarbonyl, C1-5 acyloxy, C1-5 acylamino, each of the aforementioned are optionally partially or fully halogenated, or R^b is chosen from C1-5 alkylsulphonylamino, hydroxy, oxo, halogen, nitro and nitrile;

or the pharmaceutically acceptable salts, acids or isomers thereof.

Claim 2 (currently amended): The compound according to claim 1 and wherein:

Y is $-O_{-}$, -NH-, -N(CH₂CH₃)- or -N(CH₃)-;

X is $-N(\mathbf{R}^{\mathbf{a}})$ -, or -O-;

Q is CH;

J is chosen from C1-10 alkyl, aryl or C3-7 cycloalkyl each optionally substituted by Rb;

R₂ is independently chosen from C1-6 alkyl, C3-6 cycloalkyl optionally substituted by C1-3 alkyl, acetyl, aroyl, C1-5 alkoxy, each being optionally partially or fully halogenated, halogen, methoxycarbonyl, phenylsulfonyl and -SO₂-CF₃:

each R³, R⁴ and R⁵ are hydrogen;

R^b is chosen from hydrogen, C1-5 alkyl, C2-5 alkenyl, C2-5 alkynyl, C3-8 cycloalkylC0-2 alkyl, aryl, C1-5 alkoxy, C1-5 alkylthio, amino, C1-5 alkylamino, C1-5 dialkylamino, C1-5 acyl, C1-5 alkoxycarbonyl, C1-5 acyloxy, C1-5 acylamino, C1-5 sulphonylamino, hydroxy, halogen, trifluoromethyl, nitro 3 and nitrile

or R^b is chosen from; heterocycle chosen from pyrrolidinyl, pyrrolinyl, morpholinyl, thiomorpholinyl, thiomorpholinyl sulfoxide, thiomorpholinyl sulfone, dioxalanyl, piperidinyl, piperazinyl, tetrahydrofuranyl, tetrahydrofuranyl, tetrahydrofuranyl, 1,3

dioxolanone, 1,3-dioxanone, 1,4-dioxanyl, piperidinonyl, tetrahydropyrimidonyl, pentamethylene sulfide, pentamethylene sulfoxide, pentamethylene sulfone, tetramethylene sulfoxide and tetramethylene sulfone and heteroaryl ehosen from aziridinyl, thienyl, furanyl, isoxazolyl, oxazolyl, thiazolyl, thiadiazolyl, tetrazolyl, pyrazolyl, pyrrolyl, imidazolyl, pyridinyl, pyrimidinyl, pyrazinyl, pyridazinyl, pyranyl, quinoxalinyl, indolyl, benzimidazolyl, benzoxazolyl, benzothiazolyl, benzothienyl, quinolinyl, quinazolinyl, naphthyridinyl, indazolyl, triazolyl, pyrazolo[3,4-b]pyrimidinyl, pyrrolo[2,3-b]pyridinyl, pyrazolo[3,4-b]pyridinyl, oxazo[4,5-b]pyridinyl and imidazo[4,5-b]pyridinyl.

Claim 3 (currently amended): The compound according to claim 2 and wherein:

 Ar^1 is chosen from phenyl, naphthyl, tetrahydronaphthyl, indanyl and indenyl, each Ar^1 is optionally substituted with one R^1 , and independently substituted with two R^2 groups;

Y is
$$-O_{-}$$
, $-S_{-}$ or $-N(CH_3)$ -;

 ${\bf R}^6$ is present, and is chosen from a bond, -O-, -O-(CH₂)₁₋₅-, -NH-, -C(O)-NH-, C₁₋₅ alkyl branched or unbranched, C₂₋₅ alkenyl, C₁₋₃ alkyl(OH) , heterocycle selected from morpholinyl, piperazinyl, piperidinyl, pyrrolidinyl and tetrahydrofuranyl, or aryl chosen from phenyl and naphthyl, each alkyl, alkenyl , heterocycle and aryl are optionally substituted by one to three hydroxy, C₁₋₃ alkyl, C₁₋₃ alkoxy, mono or diC₁₋₃ alkyl amino, amino or C₁₋₅ alkoxycarbonyl;

wherein each **R**₆ is further optionally covalently attached to groups chosen from: hydrogen, -N**R**₇**R**₈, C₁₋₃ alkyl, C₃₋₆ cycloalkylC₀₋₂alkyl, hydroxy, C₁₋₃ alkoxy, phenoxy, benzyloxy, phenylC₀₋₄ alkyl, piperazinylC₀₋₄ alkyl, piperidinylC₀₋₄ alkyl, pyrrolidinylC₀₋₄ alkyl, morpholinylC₀₋₄ alkyl, tetrahydrofuranylC₀₋₄ alkyl, triazolyl C₀₋₄ alkyl, imidazolyl C₀₋₄ alkyl and pyridinyl C₀₋₄ alkyl, each abovelisted heterocycle, heteroaryl and phenyl

group is optionally substituted by one to three hydroxy, oxo, C_{1-4} alkyl, C_{1-3} alkoxy, C_{1-5} alkoxycarbonyl, $-NR_7R_8$, NR_7R_8 -C(O)- or C_{1-4} acyl;

each \mathbf{R}_7 and \mathbf{R}_8 are independently hydrogen, phenyl C_{0-3} alkyl optionally subtituted by halogen, C_{1-3} alkyl or di C_{1-5} alkyl amino, or \mathbf{R}_7 and \mathbf{R}_8 are C_{1-2} acyl, benzoyl or C_{1-5} branched or unbranched alkyl optionally substituted by C_{1-4} alkoxy, hydroxy or mono or di C_{1-3} alkyl amino.

Claim 4 (currently amended): The compound according to claim 3 and wherein:

X is -O-;

 $Y is -N(CH_3)$;

J is C1-10 alkyl optionally substituted by R^b;

R₂ is independently chosen from C1-6 alkyl, C3-6 cycloalkyl optionally substituted by C1-3 alkyl and C1-5 alkoxy, each being optionally be partially or fully halogenated;

R⁶ is chosen from

a bond, -O-, -O-(CH₂)₁₋₅-, -NH-, -C(O)-NH-, C_{1-5} alkyl branched or unbranched, C_{2-5} alkenyl, C_{1-3} alkyl(OH), heterocycle selected from morpholinyl, piperazinyl, piperidinyl and pyrrolidinyl or and phenyl, each alkyl, alkenyl, heterocycle and phenyl are is optionally substituted by one to three hydroxy, C_{1-3} alkyl, C_{1-3} alkoxy, mono or diC_{1-3} alkyl amino, amino or C_{1-5} alkoxycarbonyl;

wherein each R_6 is further optionally covalently attached to groups chosen from:

hydrogen, -NR₇R₈, C₁₋₃ alkyl, C₃₋₆ cycloalkylC₀₋₂alkyl, benzyloxy, phenylC₀₋₄ alkyl, piperazinylC₀₋₄ alkyl, piperidinylC₀₋₄ alkyl, pyrrolidinylC₀₋₄ alkyl, morpholinylC₀₋₄ alkyl, triazolyl C₀₋₄alkyl, imidazolyl C₀₋₄alkyl and pyridinyl-C₀₋₄alkyl, each above-listed heterocycle, heteroaryl and phenyl group is optionally substituted by one to three

hydroxy, oxo, C_{1-4} alkyl, C_{1-3} alkoxy, C_{1-5} alkoxycarbonyl, amino, NR_7R_8 -C(O)- or C_{1-4} acyl;

each \mathbf{R}_7 and \mathbf{R}_8 are independently hydrogen, phenyl $\mathbf{C}_{0\text{-}2}$ alkyl optionally substituted subtituted by halogen, $\mathbf{C}_{1\text{-}3}$ alkyl or di $\mathbf{C}_{1\text{-}5}$ alkyl amino, or \mathbf{R}_7 and \mathbf{R}_8 are $\mathbf{C}_{1\text{-}5}$ branched or unbranched alkyl optionally substituted by $\mathbf{C}_{1\text{-}4}$ alkoxy, hydroxy or mono or di $\mathbf{C}_{1\text{-}3}$ alkyl amino;

R^b is chosen from hydrogen, C1-5 alkyl, C3-7 cycloalkylC0-2 alkyl, aryl, C1-5 alkoxy, amino, C1-5 alkylamino, C1-3 dialkylamino, C1-3 acyl, C1-5 alkoxycarbonyl, C1-3 acyloxy, C1-3 acylamino, C1-3 sulphonylamino, hydroxy, halogen, trifluoromethyl, nitro₁ and nitrile;

or R^b is chosen from pyrrolidinyl, pyrrolinyl, morpholinyl, thiomorpholinyl, thiomorpholinyl, thiomorpholinyl sulfone, piperidinyl, piperazinyl, piperidinonyl, tetrahydropyrimidonyl, aziridinyl, isoxazolyl, oxazolyl, thiazolyl, thiadiazolyl, pyrazolyl, pyrrolyl, imidazolyl, pyridinyl, pyrimidinyl, pyrazinyl and pyridazinyl.

Claim 5 (currently amended): The compound according to claim 4 and wherein:

Ar¹ is formula (A) or (B)

wherein:

when Ar^1 is formula (A) then:

 \mathbf{R}^1 is NH_2 , $\mathbf{J}\text{-}\mathrm{N}(\mathbf{R}^a)\text{-}(\mathrm{CH}_2)_{\mathfrak{m}^-}$, $\mathrm{NH}_2\mathrm{C}(\mathrm{O})\text{-}$, $\mathbf{J}\text{-}\mathrm{N}(\mathbf{R}^a)\text{-}\mathrm{C}(\mathrm{O})\text{-}$, $\mathbf{J}\text{-}\mathrm{S}(\mathrm{O})_2\text{-}\mathrm{N}(\mathbf{R}^a)\text{-}\frac{\mathrm{or}}{\mathrm{O}}$ $\mathbf{J}\text{-}\mathrm{N}(\mathbf{R}^a)$ $\mathbf{S}(\mathrm{O})_2\text{-}$ or heterocycle (CH_2)₁₋₂ wherein the heterocycle is chosen from pyrrolidinyl, morpholinyl and piperazinyl each optionally substituted by $\mathbf{C}1$ 4.alkyl, and \mathbf{J} is $\mathrm{C}_{1\text{-}5}$ alkyl optionally substituted by \mathbf{R}^b ; or when \mathbf{Ar}^1 is formula (B) then: \mathbf{R}^1 is hydrogen or halogen;

R₂ is independently chosen from C1-5 alkyl, C3-6 cycloalkyl optionally substituted by C1-3 alkyl and C1-5 alkoxy, each being optionally partially or fully halogenated;

 R^6 is chosen from a bond, -O-, -O-(CH₂)₁₋₅-, -NH-, -C(O)-NH-, C₁₋₅ alkyl branched or unbranched, C₂₋₅ alkenyl, C₁₋₃ alkyl(OH), heterocycle selected from morpholinyl, piperazinyl, piperidinyl and pyrrolidinyl or and phenyl, each alkyl, alkenyl, heterocycle and phenyl are optionally substituted by one to three hydroxy, C₁₋₃ alkyl, C₁₋₃ alkoxy, mono or diC₁₋₃ alkyl amino, amino or C₁₋₅ alkoxycarbonyl;

wherein each R₆ is further optionally covalently attached to groups chosen from:

hydrogen, -NR₇R₈, C₁₋₃ alkyl, C₃₋₆ cycloalkylC₀₋₂alkyl, benzyloxy, phenylC₀₋₄ alkyl, piperazinyl, piperazinylC₁₋₂alkyl, piperidinyl, piperidinyl C₁₋₂alkyl, pyrrolidinyl, pyrrolidinyl C₁₋₂alkyl, triazolyl C₁₋₂alkyl, imidazolyl, imidazolyl C₁₋₂alkyl, pyridinyl and pyridinyl C₁₋₂alkyl, each above-listed heterocycle, heteroaryl and phenyl group is optionally substituted by one to three hydroxy, oxo, C₁₋₄ alkyl, C₁₋₃ alkoxy, C₁₋₅ alkoxycarbonyl, amino, NR₇R₈-C(O)- or C₁₋₄ acyl.

Claim 6 (currently amended): The compound according to claim 5 and wherein: Ar^1 is formula (A) or (B)

$$\mathbb{R}^1$$
 $O-CH_3$
 $O-CH_3$
 $O-CH_3$
 $O-CH_3$

and R2 is chosen from

when Ar^1 is formula (A) then: when R^1 is J-S(O)₂- N(R^a)- or J-N(R^a)-S(O)₂- then J is C_{1-3} alkyl; and when R^1 is NH₂, J-N(R^a)-(CH₂)_m-, NH₂C(O)-, J-N(R^a)-C(O)-, or heterocycle (CH₂)₁₋₂- wherein the heterocycle is chosen from pyrrolidinyl, morpholinyl, piperazinyl or — C1-4alkylpiperazinyl, then J is C1-3 alkyl optionally substituted by R^b .

Claim 7 (currently amended): The compound according to claim 6 and wherein:

R^b is chosen from hydrogen, C1-5 alkyl, C3-6 cycloalkylC0-2 alkyl, phenyl, C1-5 alkoxy, amino, C1-5 alkylamino, C1-3 dialkylamino, C1-3 acyl, C1-5 alkoxycarbonyl, C1-3 acyloxy, C1-3 acylamino, hydroxyl; and halogen;

or R^b-is chosen from morpholinyl, thiomorpholinyl, thiomorpholinyl sulfoxide, thiomorpholinyl sulfone, piperidinyl, piperidinonyl, pyridinyl, pyrimidinyl, pyrazinyl and pyridazinyl.

Claim 8 (currently amended): The compound according to claim 7 and wherein:

R^b is chosen from amino, C1-5 alkylamino, and C1-3 dialkylamino; or R^b is chosen morpholinyl, piperidinyl and pyridinyl.

Claim 9 (original): The compound according to claim 6 and wherein: Ar^1 is formula (A).

Claim 10 (original): The compound according to claim 6 and wherein: Ar^1 is formula (B).

Claim 11 (original): The compound according to claim 6 and wherein: Ar^{1} is

$$\mathsf{CH_3S(O)_2-N} \overset{\mathsf{N}}{\underset{\mathsf{CH_3}}{\overset{\mathsf{N}}{\bigcap}}}$$

Claim 12 (currently amended): A compound chosen from:

1-Methyl-7-(pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide

7-(2-Methylamino-pyrimidin-4-yloxy)-1H-indole-2-earboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide

- 1-Methyl-7 (2-methylamino-pyrimidin-4-yloxy)-1H indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 1-Methyl-7-(2-methylamino-pyrimidin-4-yloxy)-1H indole-2-carboxylic acid (5-tert-butyl-2-methoxy-phenyl)-amide
- 7-[2-(2-Dimethylamino-ethylamino)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 7-(2-Cyclopropylamino-pyrimidin 4 yloxy) 1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 7-[2-(4-Methoxy benzylamino) pyrimidin 4-yloxy] 1-methyl-1H indole 2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino 2-methoxy phenyl) amide
- 1-Methyl-7-[2-(4-methyl-piperazin-1-yl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy phenyl) amide
- 7-[2 (2-Dimethylamino-ethylamino)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-2-methoxy-phenyl)-amide
- 1-Methyl-7-[2-(4-methyl-piperazin-1-yl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-2-methoxy-phenyl) amide
- 1-Methyl-7 [2-(2-morpholin 4-yl-ethylamino) pyrimidin 4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 7-[2-(3-Dimethylamino propylamino) pyrimidin 4-yloxy] 1 methyl 1H-indole 2-earboxylic acid (5-tert-butyl 3-methanesulfonylamino 2-methoxy phenyl) amide
- 7-[2-(3-Dimethylamino-2,2-dimethyl-propylamino) pyrimidin 4-yloxy] 1-methyl-1H-indole 2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 7-(2-Dimethylamino-pyrimidin-4-yloxy) 1-methyl-1H indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 1-Methyl-7 (6-methyl-2-methylamino-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 1 Methyl 7 [2 (2-pyrrolidin-1-yl ethylamino) pyrimidin 4 yloxy] 1H indole 2-carboxylic acid (5 tert butyl 3-methanesulfonylamino 2 methoxy phenyl) amide
- 1-Methyl-7-[2-(piperidin-4-ylamino) pyrimidin-4-yloxy] 1H indole 2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 4-{4-[2-(5-tert-Butyl-3-methanesulfonylamino-2-methoxy-phenylearbamoyl)-1-methyl-1H-indol-7-yloxy]-pyrimidin-2-ylamino}-piperidine-1-earboxylic-acid-tert-butyl-ester

- 7-{2-[(2-Dimethylamino-ethyl)-methyl-amino]-pyrimidin-4-yloxy}-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 1-Methyl-7-[6-methyl-2-(4-methyl-piperazin-1-yl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 7-[2-(2-Dimethylamino-ethoxy)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 7 [2 (2 Dimethylamino ethoxy) pyrimidin 4-yloxy] 1 methyl 1H indole 2 carboxylic acid (5 tert butyl 2 methoxy phenyl) amide
- 1-Methyl 7 [2 (2 pyrrolidin 1 yl ethoxy) pyrimidin 4 yloxy] 1H-indole 2 carboxylic acid (5 tert-butyl 3 methanesulfonylamino 2 methoxy phenyl) amide
- 1-Methyl-7-[2-(2-morpholin-4-yl-ethoxy) pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 1-Methyl-7-[2-(1-methyl-piperidin-4-yloxy)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 7-[2 (2-Dimethylamino-ethoxy)-6-methyl-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 1-Methyl-7-(2-methylcarbamoyl-pyridin-4-yloxy)-1H-indole-2-carboxylic acid [5-tert-butyl-3-(2-dimethylamino-ethylcarbamoyl)-2-methoxy-phenyl]-amide
- 7-[2-(2-Dimethylamino-ethylcarbamoyl)-pyridin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 1-Methyl-7-(2-methylamino-pyrimidin-4-yloxy)-1H indole-2-carboxylic acid [5-tert-butyl-3-(2-dimethylamino-ethylcarbamoyl) 2-methoxy-phenyl]-amide
- 1-Methyl-7-(2-methylamino-pyrimidin-4-yloxy) 1H-indole-2-carboxylic acid [5-tert-butyl-2-methoxy-3-(2-morpholin-4-yl-ethylcarbamoyl) phenyl]-amide
- 1-Methyl-7-(2-methylamino-pyrimidin-4-yloxy) 1H-indole-2-carboxylic acid (5-tert-butyl-3-carbamoyl-2-methoxy-phenyl) amide
- 1-Methyl-7-(2-methylamino-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-2-methoxy-3-methylcarbamoyl-phenyl) amide
- 1-Methyl-7 (2-vinyl-pyrimidin-4-yloxy)-1H-indole-2-earboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 7-[2-(1,2-Dihydroxy-ethyl) pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide

- 1-Methyl-7-[2-(morpholin 4-ylamino) pyrimidin 4-yloxy]-1H-indole-2-earboxylic acid (5-tert-butyl-3-methanesulfonylamino 2-methoxy-phenyl) amide
- 1-Methyl-7-(2-morpholin-4-ylmethyl-pyrimidin-4-yloxy) 1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 1-Methyl-7-(2-morpholin-4-ylmethyl-pyridin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 1 Methyl 7 [2 (4 methyl piperazin 1 ylmethyl) pyrimidin 4 yloxy] 1H indole 2 carboxylic acid (5 tert butyl 3 methanesulfonylamino 2 methoxy phenyl) amide
- 7-(2-Dimethylaminomethyl-pyridin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 1-Methyl-7-(2-methylcarbamoyl-pyridin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 7-(2-Benzyloxymethyl-pyridin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 1-Methyl-7-(2-methylamino-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-2-methoxy-3-morpholin-4-ylmethyl-phenyl) amide
- 1-Methyl-7 (2-methylamino-pyrimidin 4-yloxy) 1H indole 2-carboxylic acid [5-tert-butyl-2-methoxy-3 (4-methyl-piperazin-1-ylmethyl) phenyl] amide
- 1-Methyl-7 (2-methylamino-pyrimidin 4-yloxy) 1H indole 2-carboxylic acid (5-tert-butyl-3-dimethylaminomethyl-2-methoxy-phenyl) amide
- 1-Methyl-7-(2-methylamino-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (3-amino-5-tert-butyl-2-methoxy-phenyl) amide
- 1-Methyl-7 (2-methylamino-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-dibenzylamino-2-methoxy-phenyl)-amide
- 1-Methyl-7-(2-methylamino-pyrimidin 4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-2-methoxy-3-methylsulfamoyl-phenyl)-amide
- 7-[2 (2-Dimethylamino ethylamino) pyrimidin 4-yloxy]-1 methyl-1H-indole 2-carboxylic acid (5-tert-butyl-3-[1,3]dioxolan-2-yl-2-methoxy-phenyl) amide
- 1-Methyl-7-[2-(4-methyl-piperazin-1-yl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-2-methoxy-3-methylaminomethyl-phenyl)-amide
- 1-Methyl-7-[2-(4-methyl-piperazin-1-yl)-pyrimidin-4-yloxy]-1H-indole-2-earboxylic acid (5-tert-butyl-2-methoxy-3-pyrrolidin-1-ylmethyl-phenyl) amide

- 1-Methyl-7-{2 [methyl-(1-methyl-piperidin-4-yl)-amino]-pyrimidin-4-yloxy}-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 7-(2-Hydroxymethyl-pyridin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide

and

- 1-Methyl-7 (2-methylamino-pyrimidin 4-yloxy) 1H indole 2-carboxylic acid [5-tert-butyl-2-methoxy-3 (2-morpholin 4-yl-ethylamino) phenyl] amide
- 1-Methyl-7-(pyridin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 1-Methyl-7-(2-piperazin-1-yl-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 1-Methyl-7 (pyrimidin-4-yloxy)-1H-indole-2 carboxylic acid [3-methanesulfonylamino-2-methoxy-5 (1-methyl-cyclopropyl)-phenyl]-amide
- 1-Methyl-7-[2-(5-methyl-2,5-diaza-bicyclo[2.2.1]hept-2-yl) pyrimidin 4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 7-[2-(2,5-Diaza-bicyclo[2.2.1]hept-2-yl)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 7 (2-Methoxy pyrimidin 4-yloxy) 1-methyl 1H-indole 2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino 2-methoxy phenyl) amide
- 7-[2 (4-tert-Butyl-piperazin-1-yl) pyrimidin 4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 1-Methyl-7-[2-(2-morpholin-4-yl-ethyl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 1-Methyl-7-{2-[2-(4-methyl-piperazin-1-yl)-ethyl]-pyrimidin-4-yloxy}-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 1-Methyl-7-[2-(2-pyrrolidin-1-yl-ethyl) pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 7-[2-(2-Dimethylamino-ethyl) pyrimidin 4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide

- 1-Methyl-7-[2-(2-morpholin-4-yl-ethyl)-pyrimidin-4-yloxy]-1H-indole-2-earboxylic acid (3-methanesulfonylamino-2-methoxy-5-trifluoromethyl-phenyl) amide
- 1-Methyl-7-[2-(4-methyl-piperazin-1-yl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (3-methanesulfonylamino-2-methoxy-5-trifluoromethyl-phenyl) amide
- 7-{2-[2-(4-tert-Butyl-piperazin-1-yl)-ethyl] pyrimidin-4-yloxy}-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phonyl) amide
- 7-[2-(4-tert-Butyl-piperazin-1-ylmethyl)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 1-Methyl-7-(2-pyrrolidin-1-ylmethyl-pyrimidin 4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 7-(2,6-Dimethyl-pyridin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide and
- 7-(2-Ethyl-pyridin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 1 Methyl 7 [2-(1,2,3,6 tetrahydro pyridin-4-yl) pyrimidin 4 yloxy] 1H-indole 2-carboxylic acid (5 tert-butyl 3 methanesulfonylamino 2 methoxy phenyl) amide
- 7-(2-Amino-pyrimidin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 1-Methyl-7 (2-pyrrolidin-1-ylmethyl-pyridin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 1-Methyl-7 (2 piperidin 1 ylmethyl-pyridin 4 yloxy) 1H indole 2 carboxylic acid (5-tert-butyl 3 methanesulfonylamino 2 methoxy phenyl) amide
- 1-Methyl-7-[2-(4-methyl-piperazin-1-ylmethyl) pyridin-4-yloxy] 1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 1-Methyl-7-(pyridin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-{[(2-dimethylamino-ethyl)-methyl-amino]-methyl}-2-methoxy-phenyl)-amide
- 7-(2-{[(2-Dimethylamino-ethyl)-methyl-amino]-methyl}-pyrimidin-4-yloxy)-1-methyl-1H-indole-2-earboxylie acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 1-Methyl-7 [2-(4-methyl-piperazin-1-yl)-pyrimidin-4-yloxy]-1H-indole-2-carboxylie acid (5-tert-butyl-3-carbamoyl-2-methoxy-phenyl)-amide

- 1-Methyl-7-[2-((1S,4S)-5-methyl-2,5-diaza-bicyclo[2.2.1]hept-2-yl)-pyridin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)amide
- 1-Methyl-7-[2-(4-methyl-[1,4]diazepan-1-yl) pyridin 4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 7-(2-[1,4]Diazepan-1-yl-pyridin-4-yloxy) 1-methyl-1-H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 1-Methyl-7-(2-piperazin-1-yl-pyridin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 1-trideuterio-7-(2-piperazin-1-yl-pyridin-4-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 7-[2 (Hexahydro-pyrrolo[1,2-a]pyrazin-2-yl)-pyridin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 1-trideuterio 7-[2-(4-methyl-piperazin 1-yl) pyridin-4-yloxy]-1H indole-2-carboxylic acid [3-methanesulfonylamino-2-methoxy-5-(1-methyl-cyclopropyl) phenyl] amide
- 7-[2-((S)-3-Dimethylamino-pyrrolidin-1-yl) pyridin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid [3-methanesulfonylamino-2-methoxy-5-(1-methyl-cyclopropyl)-phenyl]-amide
- 7-[2-((S) 3-Dimethylamino-pyrrolidin-1-yl)-pyridin-4-yloxy]-1-methyl-1H-indole-2-earboxylie acid [3-methanesulfonylamino-2-methoxy-5-(1-methyl-cyclopropyl)-phenyl]-amide
- 1-Methyl-7-[2-(4-methyl-piperazine-1-carbonyl) pyridin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide and
- 1-Methyl-7 [2 (piperazine-1 carbonyl) pyridin-4-yloxy] 1H indole 2 carboxylic acid (5-tert-butyl-3 methanesulfonylamino-2 methoxy-phenyl) amide

or the pharmaceutically acceptable salts, acids or isomers thereof.

Claim 13 (currently amened): A compound chosen from:

7-(Pyrimidin-4-yloxy)-benzo[b]thiophene 2-earboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide

- 7-(Pyrimidin-4-yloxy) 1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 7-(Pyrimidin 4-yloxy)-benzofuran-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino 2-methoxy-phenyl) amide
- 1-Methyl-7-(pyrimidin-4-ylsulfanyl)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 1-Methyl-7-(pyrimidin-4-ylamino)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 1-Methyl-7-(pyridin-3-yloxy)-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 7 (2-Benzylamino-pyrimidin-4-yloxy)-1-methyl-1H-indole 2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 1-Methyl 7-{2 [(pyridin-2-ylmethyl)-amino] pyrimidin 4-yloxy}-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 7-[2-(2-Imidazol-1-yl-ethylamino) pyrimidin 4-yloxy] 1-methyl-1H-indole 2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino 2-methoxy-phenyl) amide
- 1-Methyl-7-[2-(2-[1,2,3]triazol-1-yl-ethylamino) pyrimidin 4-yloxy]-1H-indole-2-earboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 7-[2-(3-Dimethylamino-propylamino) pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid [2-methoxy-5-(2,2,2-trifluoro-1-trifluoromethyl-ethyl) phenyl]-amide
- 7-{2-[(2-Dimethylamino-ethyl)-methyl-amino]-pyrimidin-4-yloxy}-1-methyl-1H-indole-2-carboxylic acid (4-chloro-2-methoxy-5-trifluoromethyl-phenyl) amide
- 7-[2-(4-Acetyl-piperazin-1-yl) pyrimidin 4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 1-Methyl-7-[2-(4-methyl-piperazin-1-yl) pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (4-chloro-2-methoxy-5-trifluoromethyl-phenyl) amide
- 7-[2-(2-Dimethylamino-ethylamino) pyrimidin 4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (2-methoxy-5-trifluoromethoxy phenyl) amide
- 7-[2-(4-Dimethylamino-piperidin-1-yl) pyrimidin 4-yloxy] 1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide

- 7-[2-(3-Dimethylamino-pyrrolidin-1-yl)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 1-Methyl-7-[2-(1-methyl-piperidin-4-ylamino) pyrimidin-4-yloxy] 1H-indole 2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 7-[2-(1-Acetyl-piperidin-4-ylamino)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 1-Methyl-7-[2-(2-morpholin-4-yl-ethoxy)-pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid (3-methanesulfonylamino-2-methoxy-5-trifluoromethyl-phenyl) amide
- 7-[2-(2-Imidazol-1-yl-ethoxy)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-earboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl) amide
- 7-[2-(2-Imidazol-1-yl-ethoxy)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (4-chloro-2-methoxy-5-trifluoromethyl-phenyl) amide
- 7-[2 (2 Dimethylamino ethylamino) pyrimidin 4-yloxy]-1 methyl-1H-indole-2-carboxylic acid (5-tert-butyl-2 methoxy 3-methylcarbamoyl-phenyl) amide
- 7 (2-Amino pyrimidin 4-yloxy) 1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-carbamoyl-2-methoxy-phenyl) amide
- 7-(2-Amino-pyrimidin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid [5-tert-butyl-3-(2-dimethylamino-ethylcarbamoyl)-2-methoxy-phenyl]-amide
- 7-[2 (2-Dimethylamino-ethylamino)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-dimethylaminomethyl-2-methoxy-phenyl) amide
- 7-[2-(2-Dimethylamino-ethylamino) pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-2-methoxy-3-pyrrolidin-1-ylmethyl-phenyl) amide
- 7-[2 (2 Dimethylamino ethylamino) pyrimidin 4-yloxy] 1-methyl 1H-indole 2-carboxylic acid (5-tert-butyl 2-methoxy 3-morpholin 4-ylmethyl-phenyl) amide
- 1-Methyl-7 (2-morpholin-4-ylmethyl-pyrimidin-4-yloxy)-1H-indole-2-carboxylic acid (4-chloro-2-methoxy-5-trifluoromethyl-phenyl) amide
- 7-[2-(3-Dimethylamino-pyrrolidin-1-ylmethyl)-pyrimidin-4-yloxy]-1-methyl-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide
- 7-(2-Carbamoyl pyrimidin 4-yloxy) 1-methyl-1H-indole 2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino 2-methoxy phenyl) amide

- 1-Methyl-7 (2-morpholin 4-ylmethyl-pyrimidin 4-yloxy)-1H-indole-2-carboxylic acid (2-methoxy-3-morpholin 4-ylmethyl-5-trifluoromethyl-phenyl) amide
- 1-Methyl 7 [2 (4 methyl-piperazin-1 yl) pyrimidin 4 yloxy] 1H-indole 2 carboxylic acid (2 methoxy 3 morpholin 4 ylmethyl-5 trifluoromethyl-phenyl) amide
- 1-Methyl-7 (2-morpholin-4-ylmethyl-pyrimidin-4-yloxy) 1H-indole-2-carboxylic acid (3-methanesulfonylamino-2-methoxy-5-trifluoromethyl-phenyl) amide
- 7-(1'-tert-Butyl-1',2',3',4',5',6'-hexahydro-[2,4']bipyridinyl-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (3-methanesulfonylamino-2-methoxy-5-trifluoromethyl-phenyl) amide
- 1-Methyl-7-(2-methylaminomethyl-pyridin-4-yloxy)-1H-indole-2-carboxylic acid (3-methanesulfonylamino-2-methoxy-5-trifluoromethyl-phenyl)-amide
- 1-Methyl-7 (2-pyrrolidin-1-ylmethyl-pyridin-4-yloxy) 1H-indole-2 carboxylic acid (2-methoxy 3-morpholin-4-ylmethyl-5 trifluoromethyl-phenyl) amide
- 1-Methyl-7-[2-(2-morpholin-4-yl-ethyl) pyrimidin-4-yloxy]-1H indole 2-carboxylic acid (3-dimethylaminomethyl-2-methoxy-5-trifluoromethyl-phenyl) amide
- 1 Methyl 7 (2-pyrrolidin 1-ylmethyl-pyrimidin 4-yloxy) 1H indole 2-carboxylic acid (2-methoxy 3-pyrrolidin 1-ylmethyl-5-trifluoromethyl-phenyl) amide
- 7-(2-Dimethylaminomethyl-pyrimidin-4-yloxy)-1-methyl-1H indole-2-carboxylic acid [2-methoxy-3-(4-methyl-piperazin-1-ylmethyl)-5-trifluoromethyl-phenyl] amide

and

- 7-(2-Dimethylaminomethyl-pyridin-4-yloxy)-1-methyl-1H-indole-2-carboxylic acid (3-methanesulfonylamino-2-methoxy-5-trifluoromethyl-phenyl)-amide
- 7-(2-Dimethylaminomethyl-pyrimidin-4-yloxy) 1 methyl-1H indole-2 carboxylic acid (3-methanesulfonylamino-2 methoxy-5 trifluoromethyl-phenyl)-amide
- 1-Methyl-7 [2-(4-methyl-piperazin-1-ylmethyl) pyrimidin 4-yloxy] 1H-indole-2-carboxylic acid (3-methanesulfonylamino 2-methoxy-5-trifluoromethyl-phenyl) amide
- 7 (2-Dimethylaminomethyl-pyrimidin 4-yloxy) 1-methyl-1H indole 2-carboxylic acid (3-methanesulfonylamino-2-methoxy-5-trifluoromethyl-phenyl) amide
- 1-Methyl-7-[2-(2-morpholin-4-yl-ethyl) pyrimidin-4-yloxy]-1H-indole-2-carboxylic acid [3-methanesulfonylamino-2-methoxy-5-(1-methyl-cyclopropyl)-phenyl]-amide
- 1-Methyl-7-[2-(1-methyl-piperidin-4-yl) pyrimidin 4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy-phenyl)-amide

7-[2 (1-Cyclopropyl-piperidin-4-yl) pyrimidin 4-yloxy] 1-methyl 1H-indole 2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino 2-methoxy phenyl) amide

1 Methyl 7 [2 (4 methyl-piperazin-1 yl) pyrimidin-4-yloxy] 1H-indole 2-carboxylic acid (3-dimethylaminomethyl-2 methoxy 5 trifluoromethyl-phenyl) amide and

1-Methyl-7-[2-(1-methyl-pyrrolidin-3-ylamino) pyridin-4-yloxy]-1H-indole-2-carboxylic acid (5-tert-butyl-3-methanesulfonylamino-2-methoxy phenyl) amide

or the pharmaceutically acceptable salts, acids or isomers thereof.

Claim 14 (original): A pharmaceutical composition containing a pharmaceutically effective amount of a compound according to claim 1 and one or more pharmaceutically acceptable carriers and/or adjuvants.

Claim 15 (canceled).

Claim 16 (amended): A method of treating a disease or condition chosen from osteoarthritis, atherosclerosis, contact dermatitis, bone resorption diseases, reperfusion injury, asthma, multiple sclerosis, Guillain-Barre syndrome, Crohn's disease, ulcerative colitis, psoriasis, graft versus host disease, systemic lupus erythematosus, insulindependent diabetes mellitus, rheumatoid arthritis, toxic shock syndrome, Alzheimer's disease, diabetes, inflammatory bowel diseases, acute and chronic pain, stroke, myocardial infarction alone or following thrombolytic therapy, thermal injury, adult respiratory distress syndrome (ARDS), multiple organ injury secondary to trauma, acute glomerulonephritis, dermatoses with acute inflammatory components, acute purulent meningitis, syndromes associated with hemodialysis, leukopherisis, granulocyte transfusion associated syndromes, necrotizing entrerocolitis, restenosis following percutaneous transluminal coronary angioplasty, traumatic arthritis, sepsis ; and chronic obstructive pulmonary disease and congestive heart failure, said method comprising

administering to a patient a pharmaceutically effective amount of a compound according to claim 1.

Claim 17 (canceled).

Claim 18 (original): A process of making a compound of the formula (I):

$$Ar^{1} \underset{R}{\overset{O}{\underset{A}}} \underset{R^{4}}{\overset{W}{\underset{A}}} \underset{R^{6}}{\overset{R^{6}}{\underset{A}}}$$

Ar₁, X, Y, Q, W, R³, R⁴, R⁵, R⁶ and R^y are defined in claim 1; said process comprising

coupling under suitable conditions an amine bearing Ar¹ carboxylic acid of the formula (III), where P is a protecting group,

removing the protecting group P to provide an intermediate of formula (V) under suitable conditions;

coupling under suitable conditions the intermediate (V) with a halo heterocycle VI (Z = halogen) bearing R^6 in the presence of a suitable base to provide a compound of the formula (I):